

AMENDMENTS TO THE CLAIMS

A detailed listing of all claims that are, or were, in the present application, irrespective of whether the claim(s) remain(s) under examination in the application is presented below. The claims are presented in ascending order and each includes one status identifier. Those claims not cancelled or withdrawn but amended by the current amendment utilize the following notations for amendment: 1. deleted matter is shown by strikethrough for six or more characters and double brackets for five or fewer characters; and 2. added matter is shown by underlining.

1. (Currently Amended) A catalyst for use in a Fischer-Tropsch synthesis reaction, the catalyst comprising ~~which comprises~~ cobalt supported on alumina, in which: the catalyst average particle size is in the range 20 to 100 μm ; the specific surface area of the impregnated and calcined catalyst particles is greater than 120 m^2/g ; the average pore size of the impregnated and calcined catalyst is at least 90 \AA (9nm); ~~[[and]]~~ the pore volume of the impregnated and calcined catalyst is at least 0.45 cm^3/g ; the cobalt content of the catalyst is from 10 to 40% by weight.
2. (Previously Presented) A catalyst as claimed in Claim 1, in which the specific surface area of the impregnated and calcined catalyst particles is in the range 120 to 220 m^2/g .
3. (Original) A catalyst as claimed in Claim 1, in which the particle size range is 40 to 80 μm .
4. (Previously Presented) A catalyst as claimed in Claim 1, in which the average pore size of the impregnated and calcined catalyst is at least 110 \AA (11nm).
5. (Previously Presented) A catalyst as claimed in Claim 4, in which the average pore size of the impregnated and calcined catalyst is at least 130 \AA (13 nm).
- 6-7. (Cancelled)

8. (Previously Presented) A catalyst as claimed in Claim 1, incorporating less than 3% by weight of a promoter.
9. (Original) A catalyst as claimed in Claim 8, in which the promoter is rhenium or platinum.
10. (Previously Presented) A catalyst as claimed in Claim 1, in which the support material is γ -alumina.
11. (Original) A catalyst as claimed in Claim 10, in which the γ -alumina is stabilised with a stabilising agent.
12. (Original) A catalyst as claimed in Claim 11, in which the γ -alumina is stabilised with lanthanum.
13. (Previously Presented) A catalyst as claimed in Claim 1, in which the alumina support includes a binder.
14. (Original) A catalyst as claimed in Claim 13, in which the binder represents less than 25% by weight of the catalyst.

15. (Previously Presented) A catalyst as claimed in Claim 13, in which the binder is an alumina-containing binder material.
16. (Previously Presented) A catalyst as claimed in Claim 1, in which the specific surface area of the prepared catalyst, comprising the cobalt in an active catalytic form on the support, is in the range 125 to 160 m²/g.
17. (Cancelled)
18. (Currently Amended) A catalyst as claimed in Claim 1[[7]], in which the cobalt content is from 15 to 25% by weight.
19. (Currently Amended) A process for the production of a catalyst ~~as claimed in any preceding Claim~~, which comprises: impregnating an alumina support with cobalt and optionally a promoter, optionally drying at less than 120°C, calcining the impregnated support at a temperature in the range 300 to 500°C and treating the calcined catalyst with a reducing gas at an activation temperature in the range 250 to 500°C; the alumina support prior to impregnation having a specific surface area in the range 80 to 225 m²/g and a pore diameter in the range 110 to 400Å (11 to 40nm).
20. (Withdrawn) A process as claimed in Claim 19, in which the alumina support has a pore volume in the range 0.6 to 1.0 cm³/g, prior to impregnation.

21. (Withdrawn) A process as claimed in Claim 19, in which the peak calcination temperature is in the range 300 to 450°C.
22. (Withdrawn) A process as claimed in Claim 19, in which the activation temperature is in the range 300 to 500°C.
23. (Withdrawn) A process as claimed in Claim 22, in which the activation temperature is in the range 300 to 450°C.
24. (Withdrawn) A process as claimed in Claim 19, in which the calcination is carried out for between 0.5 and 6 hours.
25. (Withdrawn) A process as claimed in Claim 19, in which the activation treatment is carried out for between 1 and 10 hours.
26. (Withdrawn) A process as claimed in Claim 19, in which the reducing gas is hydrogen and/or carbon monoxide, optionally mixed with an inert gas.
27. (Withdrawn) A process as claimed in Claim 19, in which, prior to impregnation, the support is pre-calcined at a temperature in the range of 400 to 900°C.

28. (Withdrawn) A process as claimed in Claim 19, in which the alumina support is γ -alumina and the process includes the step of stabilising the γ -alumina prior to the calcination step.
29. (Withdrawn) A process as claimed in Claim 19, in which prior to impregnation, the alumina support has a specific surface in the range 150 to 240 m²/g.
30. (Withdrawn) A process as claimed in Claim 19, in which, prior to impregnation, the alumina support has a pore volume in the range 0.7 to 0.9 cm³/g.
31. (Withdrawn) A process as claimed in Claim 19, in which the impregnation step comprises an incipient wetness treatment in which an aqueous solution of a cobalt compound and optionally a rhenium compound is mixed with the dry support material until the pores are filled, and the impregnated support is then dried, prior to the calcining step.
32. (Withdrawn) A process as claimed in Claim 31, in which the amount of aqueous solution used in the impregnation is 0.05-2 times larger than the measured pore volume of the catalyst support.
33. (Withdrawn) A process as claimed in Claim 31, in which drying is carried out at 80 to 120°C.

34. (Withdrawn) A process as claimed in Claim 31, in which the cobalt compound is selected from cobalt nitrate ($\text{Co}(\text{NO}_3)_2$), cobalt acetate(s), cobalt halide(s), cobalt carbonyl(s), cobalt oxalate(s), cobalt phosphate(s), cobalt carbonate(s), cobalt (hexa)amine salt(s) and organic cobalt compounds.
35. (Withdrawn) A process as claimed in Claim 31, in which the rhenium compound is selected from perrhenic acid (HReO_4), ammonium perrhenate, rhenium halide(s) and rhenium carbonyl(s).
36. (Withdrawn) A process as claimed in Claim 35, in which the cobalt compound is cobalt nitrate and the rhenium compound is perrhenic acid.
37. (Withdrawn) A process as claimed in Claim 19, in which, prior to impregnation, the alumina support has an ASTM attrition value of less than 30% by weight of fines produced by 5 hours testing.
38. (Withdrawn) A process as claimed in Claim 37 in which the ASTM value is less than 20%.
39. (Withdrawn) The use of a catalyst as claimed in Claim 1, in a Fischer-Tropsch synthesis reaction.

40. (Withdrawn) The use of a catalyst manufactured according to a process as claimed in Claim 19, in a Fischer-Tropsch synthesis reaction.
41. (Withdrawn) The use of a catalyst as claimed in Claim 39, in which the reaction is carried out in a slurry bubble column reactor.
42. (Withdrawn) A use as claimed in Claim 41, in which H_2 and CO are supplied to a slurry in the reactor, the slurry comprising the catalyst in suspension in a liquid including the reaction products of the H_2 and CO, the catalyst being maintained in suspension in the slurry at least partly by the motion of the gas supplied to the slurry.
43. (Withdrawn) A process for the production of hydrocarbons which comprise subjecting H_2 and CO gases to a Fischer-Tropsch synthesis reaction in a reactor in the presence of a catalyst as claimed in Claim 1.
44. (Withdrawn) A process for the production of hydrocarbons which comprise subjecting H_2 and CO gases to a Fischer-Tropsch synthesis reaction in the presence of a catalyst manufactured according to a process as claimed in Claim 19.
45. (Withdrawn) A process as claimed in Claim 43, in which the reaction is a three-phase reaction in which the reactants are gaseous, the product is at least partially liquid and the catalyst is solid.

46. (Withdrawn) A process as claimed in Claim 45, in which the reaction is carried out in a slurry bubble column reactor.

47. (Withdrawn) A process as claimed in Claim 46, in which the H₂ and CO are supplied to a slurry in the reactor, the slurry comprising the catalyst in suspension in a liquid including the reaction products of the H₂ and CO, the catalyst being maintained in suspension in the slurry at least partly by the motion of the gas supplied to the slurry.

48. (Withdrawn) A process as claimed in Claim 43, in which the reaction temperature is in the range 190-250°C.

49. (Withdrawn) A process as claimed in Claim 48, in which the reaction temperature is in the range 200-230°C.

50. (Withdrawn) A process as claimed in Claim 43, in which the reaction pressure is in the range 10-60 bar.

51. (Withdrawn) A process as claimed in Claim 50, in which the reaction pressure is in the range 15 to 30 bar.

52. (Withdrawn) A process as claimed in Claim 43, in which the H_2/CO ratio of the gases supplied to the Fischer-Tropsch synthesis reactor is in the range 1.1 to 2.2.

53. (Withdrawn) A process as claimed in Claim 52, in which the H_2/CO ratio is in the range 1.5 to 1.95.

54. (Withdrawn) A process as claimed in Claim 43, in which the superficial gas velocity in the reactor is in the range 5 to 60 cm/s.

55. (Withdrawn) A process as claimed in Claim 54 in which the superficial gas velocity is in the range 20 to 40 cm/s.

56. (Withdrawn) A process as claimed in Claim 43, in which the product of the Fischer-Tropsch synthesis reaction is subsequently subjected to post-processing.

57. (Withdrawn) A process as claimed in Claim 56 in which the post-processing is selected from de-waxing, hydro-isomerisation, hydro-cracking and combinations of these.

58-63. (Cancelled)